

LIPERT, Z. [Lippert, Z.], inzh. (Praga); MEDEK, R. (Praga)

Conic cogwheels with screw teeth, and milling heads made with the aid of universal metal-cutting tools. Mashinostroene 11 no.4:17-22 Ap '62.

LIPPICH, Iaszlo

Measuring and topographic instruments in civil engineering. Good. kart.
12 no.1:41-45 '60. (REAL 9:5)
(Civil engineering) (Topographic surveying)

LIPPICH, Laszlo

Taxation problems of works on expropriation plans. Geod kart 12 no.4:
274-278 '60. (EKAI 10:3)
(Hungary--Geodesy)

LIPPICH, Laszlo; MOLNAR, Pal; PASZTHY, Miklos

Surveying and tracing works in connection with wire-rope ways. Geod
kart 13 no.1:22-25 '61. (EEAI 10:6)
(Wire-rope transportation)

LIPPICH, Laszlo

Expropriation operations. Geod kart 13 no.2:115-119 '61.

1. LIPPING, V. O.
2. USSR (600)
3. Horse Breeding
4. "From the history of Russian horse breeding." V. O. Vitt. Reviewed by V. O. Lipping.
Konevodstvo, No. 11 - 1952.

9. Monthly List of Russian Acquisitions, Library of Congress, February, 1953. Unclassified.

LIPPMAA, E.

Use of double resonance in studying the spectra of nuclear
magnetic resonance. Izv. AN Est. SSR. Ser. fiz.-mat. i tekhn.
nauk 14 no.1:125-128 '65. (MIRA 18:11)

1. Institut kibernetiki AN Estonskoy SSSR.

LIPMAA, E.; SYUGIS, A. [Syugis, A.]

Side band spin-generator with phase-synchronized modulation frequency. Izv. AN Est. SSR. Ser. fiz.-mat. i tekhn. nauk 14 no.1:129-132 '65. (MIRA 18:11)

1. Institut kibernetiki AN Estonskoy SSR.

LIPPMAA, E.; LUYGA, P. [Luiga, P.]

Nonuniformity of gas flow in preparative gas chromatography.
Izv. AN Est. SSR. Ser.fiz.-mat.1 tekhnauk 14 no.2:246-254
'65.

Loading factor and condensation in preparative gas chromatography. Ibid.:255-257 (MIRA 19:1)

1. Institut kibernetiki AN Estonskoy SSR. Submitted June 29, 1964.

SINIVEE, V.; LIPPMAA, E.

Weak perturbing radiofrequency field effects in nuclear magnetic double resonance. Part 1. Izv.AN Est. SSR. Ser.fiz.-mat.i tekhnauk 14 no.2:258-265 '65.

(MIRA 19:1)

1. Academy of Sciences of the Estonian S.S.R., Institute of Cybernetics.

LUIGA, P.; LIPPMAA, E.

Use of the thermal pulse method in preparative gas chromatography.
Izv. AN Est. SSR. Ser.fiz.-mat.i tekhnauk 14 no.2:305-306 '65.
(MIRA 19:1)

1. Institut kibernetiki AN Estonskoy SSR. Submitted March 17,
1965.

LIFPMAN, E.; PUKHAR, Yu.; ACLA, M.; STUGIS, A. [Sugis, A.]

Use of nuclear magnetic double resonance with a weak perturbing ("tickling") field in determining the mutual disposition of energy levels in a spin system. Part 1. Izv. AN Est. SSR. Ser. fiz.-mat. i tekhn. nauk 14 no.2:306-307 '65.

(MIRA 19:1)

1. Institut kibernetiki AN Estonskoy SSR. Submitted March 17, 1965.

PAST, Ya. [Past, J.]; LIEPMAA, M.; OLIVSON, A.

Use of the modulation of the magnetic field in nuclear magnetic resonance spectrometry of carbon-13. Izv. AN Est. SSR. Ser.fiz.-mat. i tekhnauk 14 no.2:308 '65. (MIRA 19:1)

1. Institut kibernetiki AN Estonskoy SSR. Submitted March 20, 1965.

LIPPMAA, E.; OLIVSON, A.; PAST, Ya. [Past J.]

Nuclear magnetic resonance in carbon-13. Part 1. Izv. AN Est.
SSR. Ser. fiz.-mat. i tekhn. nauk 14 no.3:477-486 '65.
(MIRA 18:11)

1. Institut kibernetiki AN Estonskoy SSR.

LIPPMAA, E.; PUSKAR, Yu. [Puskar, J.]; ALLIA, M.

Use of the method of double intermolecular magnetic resonance
(INDOR) in studying the nuclear Overhauser effect. Izv. AN Est.
SSR. Ser. fiz.-mat. i tekhn. nauk 14 no.3:487-489 '65.
(MIRA 18:11)

1. Institut kibernetiki AN Estonskoy SSR.

SINIVEE, V.; LIPPMAA, E.

Effects of a weak perturbing radio-frequency field in double nuclear magnetic resonance. Part 2. Izv. AN Est. SSR. Ser. fiz.-mat. i tekhn. nauk 14 no. 4:564-568 '65 (MIRA 19:2)

1. Institut kibernetiki AN Estonskoy SSR. Submitted July 29, 1965.

LIPPING, V.O., kand. sel'skokhozyaystvennykh nauk.

"Artificial insemination of farm animals; manual for zootechnicians and veterinarians" by F.V. Ozhin and others. Reviewed by V.O. Lipping.
Zhivotnovodstvo 20 no.2:89-92 F '58. (MIRA 11:1)
(Artificial insemination)
(Ozhin, F.V. and others)

6117 A -

✓ Acidimetric determination of phenols. E. T. Lippman
(Polytech. Inst., Tallin, Estonia). *Zhur. Anal. Khim.* 10:
189-74; *J. Anal. Chem. U.S.S.R.* 10, 157-62(1965)
(Engl. translation).—Phenols, enols, and carboxylic acids
were titrated in a specially designed high-frequency con-
ductometric app. As solvent a mixt. of diethylamine
(tech., contg. approx. 30% triethylamine) 450, diethyl-
formamide 350, pyridine 200 ml., and thymol 1 g. was used.
The titrant was 0.3N K methylate prepd. from pyridine
840, benzene 80, MeOH 100 ml., and K 13 g. M. Hozch

17 80

LIPPMAA, E. T.

LIPPMAA, E. T.: "The dynamics of the isolation of products of thermal decomposition of Estonian oil shale." Min Higher Education USSR. Tallin Polytechnic Ist. Tallin, 1956. (Dissertation for the Degree of Candidate in Technical Science.)

Knizhnaya Letopis'
No 32, 1956. Moscow.

"APPROVED FOR RELEASE: 07/12/2001

CIA-RDP86-00513R000930030008-8

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100

APPROVED FOR RELEASE: 07/12/2001

CIA-RDP86-00513R000930030008-8"

LIPPMAA, E.T.

1445. STRUCTURE OF THE HYDROCARBON PORTION OF KERAGEN FROM ESTONIAN SHALE. Aarva, A.Ya, and LIPPMAA, E.T. (Inst. Khim. Akad. Nauk SSSR, Moscow), 1977, vol. 30, 449-456, 15 refs. (Engl. abstr. 1977, vol. 11, 13469). A Kerogen concentrate obtained by centrifuging a benzene solution was (a) diazotized with p-nitrobenzenediazonium sulphate and with 2,4,6-trinitrobenzenediazonium sulphate; (b) diazotized with a solution of paraformaldehyde in concentrated hydrochloric acid; and (c) diazotized with mercuric acetate. Methods a, b, and c indicated the presence of 0.16, and 0.23 equivalents of aromatic rings and 0.16 equivalents of the oxygen in kerogen was in the aromatic ring; also, the phenols were destroyed and were formed during the initial decomposition of kerogen. Paraffinic aliphatic double bonds were present; 0.16 equivalent of oxygen was in the form of carbonyls. The carbon distribution in kerogen consists, approximately, of 19% in aromatic rings, 50-75% in naphthenic rings, and 5-35% in the paraffinic chains.

C.A.

ns

LIPPMANA,
AARNA, A.Ya. [Aarna, A.J.]; LIPPMAN, E.T.; PALUOYA, V.T. [Paluoja, V.T.]

Properties of neutral oxygen compounds of shale tar. Khim. i
tekh. gor. slan. i prod. ikh perer. no.9:139-146 '60. (MIRA 15:6)

(Kivioli—Oil shales—Analysis)

L 14819-65 EWT(1)/EEC(t) Feb IJP(c)/AFWL/SSD/AS(mp)-2/RAEM(c)/RAEM(i)/ESD(t)

ACCESSION NR: AR3004149

S/0272/63/000/006/0142/0142

SOURCE: RZh. Metrologiya i izmer. tekhn. Otd. vy*p., Abs. 6.32.1120

AUTHOR: Lippmaa, E.T.

TITLE: A nuclear magnetic resonance spectrometer of high resolving power, equipped with spin stabilization

CITED SOURCE: Tr. Tallinsk. politekh. in-ta, A, no. 195, 1962, 65-77

TOPIC TAGS: nuclear magnetic resonance, magnetic resonance spectrometer, high resolution spectrometer, spin stabilization, spectrometer design

TRANSLATION: The article describes a spectrometer with spin stabilization of the ratio of frequency to magnetic field intensity, designed at the Tallinskiy politeknicheskiy institut (Tallin Polytechnical Institute). The author discusses the theoretical principles of the stabilization method, presents schematic illustrations of the entire instrument and its individual components, and discusses the operation of the spin generator. Spectra of the same materials, obtained with and without the spin stabilizer, are evaluated comparatively. Bibl. with 35 titles; 6 illustrations. M. Mekler

Card 1/2

L 14819-65

ACCESSION NR: AR3004149

SUB CODE: NP, OP

ENCL: 00

Card 2/2

L 3539-66 EWT(1)/EPF(c)/EWA(h) IJP(c) WIV/CG
 ACCESSION NR: AP5015747 UR/0023/65/000/001/0129/0132

AUTHORS: Lippmaa, E.; Sugis, A. (Syugis, A.) 44, 45 44 41 B

TITLE: Investigation of the sideband spin generator with phase synchronized modulation frequency

SOURCE: AN EstSSR. Izvestiya. Seriya fiziko-matematicheskikh i tekhnicheskikh nauk, no. 1, 1965, 129-132

TOPIC TAGS: stabilizer, spin resonance, nuclear resonance, epr spectrometry, signal generator 25

ABSTRACT: The authors describe a spin generator operating at 40 Mc, with a modulation frequency of 5 or 10 kcs, in which the modulation frequency is phase-synchronized with an external stable frequency to make it possible to use the generator for experiments with double resonance. The generator is used as a stabilizer in a nuclear double resonance spectrometer. A block diagram of the spin generator is shown in Fig. 1 of the Enclosure. The audio-frequency signal (5 or 10 kcs) from the high-frequency phase detector is clipped in a

Card 1/3

L 3539-66

ACCESSION NR: AP5015747

3
diode limiter, filtered, and applied to the modulation coils through a power amplifier. To hold the modulation frequency constant, the filtered audio signal is fed to a phase detector where it is compared with a very stable audio signal from a quartz oscillator. The output voltage is applied to a parametric diode of the high-frequency oscillator, whose frequency was made to follow exactly the variations of the magnetic field. The stabilization coefficient at 10 kcs is 1250 and increases with decreasing operating frequency; it falls to 70 per cent of its initial value at 250 cps. Orig. art. has: 1 figure and 4 formulas

ASSOCIATION: Institute kibernetiki AN ESSR (Institute of Cybernetics, AN ESSR)

SUBMITTED: 06Jan65

ENCL: 01

SUB CODE: NP

NR REF SOV: 004

OTHER: 011

Card 2/3

L 3539-66
ACCESSION NR: AP5015747

ENCLOSURE: 01

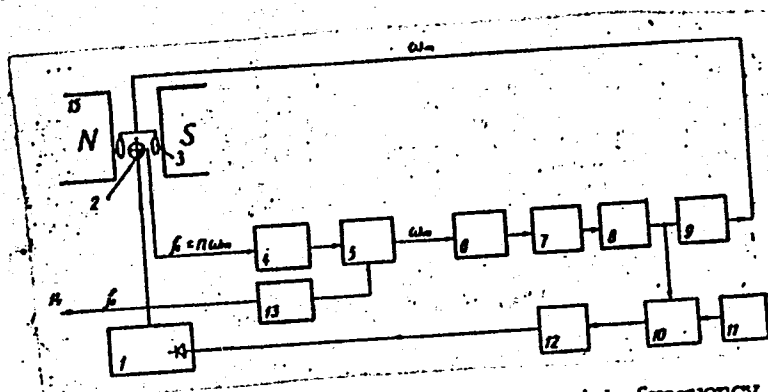


Fig. 1. Block diagram of spin generator. 1 - High-frequency generator, 2 - crossed coils, 3 - modulation coils, 4 - hf amplifier, 5 - phase detector, 6 - 1f amplifier, 7 - diode limiter, 8 - filter, 9 - input stage of 1f amplifier, 10 - phase detector, 11 - 1f quartz generator, 12 - filter, 13 - phase shifter, 14 - hf output, 15 - electromagnet.

Card 3/3

ACC NR: AT7005782

SOURCE CODE: UR/2807/66/000/238/0003/0018

AUTHORS: Kukk, P. L.; Syugis, A. Yu.; Varvas, Yu. A.; Lippmaa, E. T.

ORG: none

TITLE: Investigation of the noise spectrum of polycrystalline cadmium sulfide

SOURCE: Tallinn. Politekhnikheskiy institut. Trudy. Seriya A, no. 238, 1966. Sbornik statey po khimii i khimicheskoy tekhnologii (Collection of articles on chemistry and chemical engineering), no. 15, 3-18

TOPIC TAGS: ^{noise spectrum, radio noise,} photoelectric effect, photoresistor, photodiode, photoconductor, cadmium sulfide / S-092 photoresistor

ABSTRACT: The noise spectrum of polycrystalline cadmium sulfide photoresistor S-092 was investigated. This work supplements the results of Yu. A. Varvas and P. L. Kukk (Trudy TPI, seriya A, No. 230, str. 109, 1965). A brief literature survey of pertinent papers dealing with the theory of experimental determination of noise in CdS photoresistors is presented, and a schematic of the experimental installation is included. The experimental results are shown graphically (see Fig. 1). It was found that the noise photocurrent in the resistor S-092 may be represented by the expression

$$S_i = \text{const. } U^2 \Phi f^{-1/2}$$

Card 1/2

UDC: 621.372.001.2

ACC NR: AT7005782

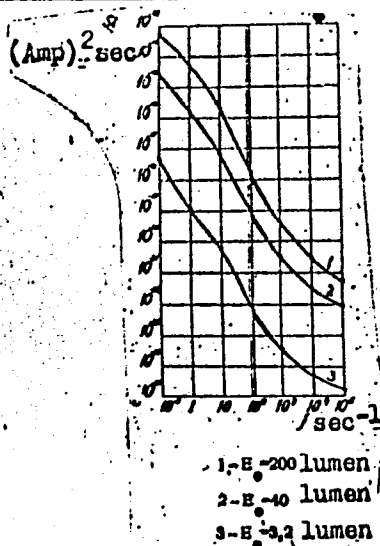


Fig. 1. Noise spectrum density S_1 as a function of the light intensity E_0 and of noise frequency f

where U is the imposed voltage, ϕ - the light density in watts/cm², and f - the frequency of the incident light. It was found that the particular position of the measuring electrodes had no noticeable effect on the shape of the noise spectrum. Orig. art. has: 8 graphs and 10 equations.

SUB CODE: 20/

SUBM DATE: none/

ORIG REF: 016/

OTH REF: 061

Card 2/2

HOMMIK, K., kand. tekhn. nauk; KALJUNAE, H., inzh. gidrotekhn.;
KASK, R., kand. sel'khoz. nauk; KATUS, A., inzh. lesnogo khoz.;
KILDEMAA, K., kand. geogr. nauk; KURKUS, J., agronom; LIPPMAA, A.,
inzh. gidrotekhn.; PANT, R., prepodavatel', agronom; RAIG, V.,
inzh. gidrotekhn.; REMMEL, A., inzh. melior.; TALPSEPP, E., kand.
sel'khoz. nauk; SOOSAAR, V., inzh., lesnogo khoz.; STERNFELD, R.,
inzh. stroit.; TOMINGAS, E., inzh. melior.; KARUS, G., red.;
RAUD, M., red.; VANTRE, I., tekhn. red.

[Handbook for soil improvement] Maaparanduse kasiraamat. Tal-
linn, Eesti riiklik kirjastus. Vol.1. [Fundamentals of soil
improvement] Maaparanduse alused. 1962. 473 p. (MIRA 15:5)
(Soils)

LIPPOCZY, B.

"What is the influence of the quantity of precipitation on acorn production?" p. 149.

AZ ERDO. (Orszagos Erdeszeti Egyesulet). Budapest, Hungary, Vol. 11, No. 6, June 1959.

Monthly list of East European Accessions (EEAI), LC, Vol. 8, No. 8, August 1959.
Uncla.

LIPPOMAN, Maria; NOWICKI, Boleslaw

Attempt to estimate the breeding value of cow families. Prace
nauk roln i lesn 12 no.4:57-74 '62 [publ. '63]

LIPPONEN, V.I., gornyy inzh.; TEOKHAROV, N.B., gornyy inzh.; TSOY, V.Ch.,
gornyy inzh.

Attachment for balancing parts. Gor. zhur. no.5:69 My '63.
(MIRA 16:5)

(Balancing of machinery)

POL

Reactions of aliphatic nitro compounds. X. Formation of the hexahydropyrimidine ring using 1-nitropropane, formaldehyde, and ammonia. Tadeusz Urbanski, Zbigniew BERNACKI, and Ewa LISAK. *Roczniki Chem.* 23, 169-71 (1954) (English summary). Cf. preceding abstr. ---PrNO_2 , CH_2O , and NH_3 in a 1:3:3 molar ratio at 95-6° for 1.5 hrs. give 3% 5-nitro-2-ethylhexahydropyrimidine (I), m. 159-60°. A 12.5% yield of I is obtained by dissolving 55 g. 2-nitro-2-ethyl-1,3-propanediol in 120 ml. 25% NH_3 , extg. with CaH_2 for several weeks and treating with alc. HCl to obtain $\text{I}\cdot\text{HCl}$, m. 150°, which reacts with 20% NaNO_2 to give the 1,3-dinitroso deriv., m. 116°. Boiling this with HCl gives the dihydrochloride of I, m. 157-8°. XI. A new derivative of tetrahydroxazine with nitromethane, formaldehyde, and benzylamine. Tadeusz Urbanski and Daniela GARG. *Ibid.* 175-81. One mole of $(\text{HOCH}_2)_2\text{CNO}_2$ and 1 mole of benzylamine with app. 3 moles of 30% CH_2O at 15° warmed for 3 hrs. at 65°, after the initial reaction subsides there is obtained 42% 5-nitro-5-hydroxymethyl-3-benzyltetrahydro-1,3-oxazine (I), m. 142-4°. To 7.5 g. I, and 4 g. NaOH in 20 H_2O is slowly added 39 ml. 30% H_2O_2 at 20°, the mixt. stirred for 1 hr., decompd. with HOAc , extd. with Et_2O , and evapd. Treatment of the oily residue with alc. HCl gives 2 g. 5-nitro-3-benzyltetrahydro-1,3-oxazine- HCl , m. 210-12°. I (4 g.) reduced with 3.5 moles H in the presence of Pd gives the 5-amino deriv. Chester Placek

MIHAILESCU, Matei; MUNTEANU, E.; LIPSCHUTZ, G.

The role of inguino-iliac adenectomies in the course and treatment of tuberculous osteoarthritis and synovitis of the lower limb.
Rumanian M. Rev. 3 no.4:25-27 O-D '59.

1. Surgical Hospital for Bone Tuberculosis, Bucharest.
(TUBERCULOSIS, OSTEOARTICULAR, surgery)
(SYNOVITIS, surgery)
(LYMPH NODES, surgery)

LIPSHITS, A.M., inzh.

Membrane-type flexible sensitive elements for power compensated differential manometer pickups. Priborostroenie no. 10: 30-33 0 '65. (MIRA 19:1)

S/119/60/000/06/14/016
B014/B014

AUTHOR: Lipshits, A. M., Engineer

TITLE: All-Union Scientific-technical Conference on the Use of
Elastic Sensitive Elements in Instrument Construction

PERIODICAL: Priborostroyeniye, 1960, No. 6, pp. 31-32

TEXT: The first All-Union Scientific-technical Conference on the Use of Elastic Sensitive Elements in Instrument Construction took place from March 22 to 25, 1960. The lectures delivered there dealt with the present stage of the development of elastic sensitive elements, its prospects in the future, and with its theory and calculation. As is known, these sensitive elements are used in pressure gauges and pneumatic control systems, and some of their advantages and disadvantages are then described. The delegates stated that the level of theoretical investigations and the methods of calculation are still insufficient. An exhibition of elastic sensitive elements took place at the same time. Particular attention was devoted to lectures dealing with the methods of producing elastic sensitive elements. The advantageous use of sheet

Card 1/2

All-Union Scientific-technical Conference on
the Use of Elastic Sensitive Elements in
Instrument Construction

S/119/60/000/06/14/016
B014/B014

materials instead of tubes for sensitive elements is stressed. A few other lectures dealt with the study and stabilization of the characteristic features of sensitive elements, and it was pointed out that new materials have been developed by the OKB and various scientific research institutes. Two new alloys for springs have been developed by the TsNIIChermet. Next, the author describes some other alloys, and points out that many new alloys are insufficiently used in industry. Finally, the main problems arising in the further development of elastic sensitive elements are summarized. These are the establishment of new design offices, supply of high-quality materials, centralization of development, standardization, specialization in the production of various sensitive elements, and the development of a usable theory. ✓

Card 2/2

L 27918-66

ACC NR: AP6017711

SOURCE CODE: UR/0119/65/000/010/0030/0032

AUTHOR: Lipshits, A. M. (Engineer)

ORG: none

TITLE: Elastic diaphragm sensing elements with force compensation for differential manometer gauges

SOURCE: Priborostroyeniye, no. 10, 1965, 30-32

TOPIC TAGS: manometer, mechanical engineering

ABSTRACT: The author studies variations in the effective area of diaphragm elements as a function of the geometric parameters of the diaphragm in diaphragm cases based on the principle of force compensation. Diaphragm cases of this type are used for protecting the measuring units in differential manometers from unilateral overloading by excess pressure and for reducing the rigidity of sensing elements. Formulas are derived for calculating the effective area of the diaphragm element as a function of profile parameters. It was found that the effective area in diaphragm cases with a serrated profile may be increased or decreased depending on the form of corrugation used. Tests showed that a minimum change in effective area with pressure variation can be assured by using corrugations 0.25 mm deep. Diaphragm cases with radical corrugations were found to be the most effective. L. Ye. Andreyeva, T. L. Lodochnikov and A. G. Shadhin participated in the work. Orig. art. has: 3 figures, 2 formulas and 2 tables. [JPRS]

SUB CODE: 13 / SUBM DATE: none / ORIG REF: 002

Card 1/1 018 UDC: 62-278:621.3.083.8

LIPSHITS, B. A.

Kashevarov, A. F.

Alaskan explorer A. F. Kashevarov. Sov. etn. no. 1, 1952.

Monthly List of Russian Accessions, Library of Congress, August 1952., Unclassified.

LIPSHITS, B.A.; TOKAREV, S.A. [reviewers]; LIPS, Julius [author].

How American imperialism corrupts the negro intelligentsia ("Journey into twilight" [in German]. Julius Lips. Reviewed by B.A.Lipshits, S.A.Tokarev).
Sov.etn. no.4:170-180 '53. (MLRA 6:12)

(Lips, Julius Ernst, 1895-1950) (United States--Negroes)
(Negroes--United States) (Howard University)

LIPSHITS, B.A.

Collections of the Museum of Anthropology and Ethnography collected
by Russian travellers and explorers in Alaska and California. Sbor.
Muz. ant. i etn. no. 16:358-369 '55. (MLRA 8:11)
(Alaska--Handicraft) (California--Handicraft)

LIPSHITS, B-M

12

JE2C

✓ Viscosity and static shear stress of alkaline melts containing antimonate or arsenate of sodium, azide reducing

of lead

H. A. Lipshits
242
chem

NaCl lowers the soly. of Na_2SiO_3 in NaOH . Hence the melts obtained by dissolving Na_2SiO_3 in NaOH with NaCl are more viscous than those obtained by dissolving Na_2SiO_3 in NaOH alone.

1.11 1.15

7
Solubility of zinc oxide in aqueous solutions of sodium hydroxide. K. G. Orlov, B. M. Lipshitz, and I. A. Lovchikov. *Tretye MZSN* 20, 700 (1972).
The soly. in the system $\text{Na}_2\text{O}-\text{H}_2\text{O}-\text{ZnO}$ was detd. at 25° and 75°. The isotherms consist of 3 branches. At 25° the

branches cover the ranges of 0-11.4, 11.4-24.70, and 24.70-39.54% Na_2O , and the tie lines of each branch intersect at points corresponding to the ratios of $\text{ZnO}:\text{H}_2\text{O}:\text{Na}_2\text{O}$ of 1:0:1, 1:0:0, and 1:5:2, resp. Each branch is in equil. with the solid phases of $\text{Zn}(\text{OH})_2$, ZnO , and $[\text{Zn}(\text{OH})_4]^{2-}\text{Na}_2\text{O}$, resp. The soly. of the first 2 phases increases with the NaOH concn. and that of the 3rd decreases. At 75° the 3 branches cover the ranges of 0-32, 32.2-44.8, and 44.8-55% Na_2O , and the tie lines of each intersect at points corresponding to ratios of $\text{ZnO}:\text{H}_2\text{O}:\text{Na}_2\text{O}$ of 1:0:0, 1:0.6:1.5, and 1:1:5. Each is in equil. with the solid phases ZnO , $[\text{Zn}(\text{OH})_4]^{2-}\text{Na}_2\text{O}$, and $[\text{Zn}(\text{OH})_4]^{2-}\text{Na}_2\text{O}$, resp. The soly. over the 2nd and 3rd branch is given by % $\text{Zn} \approx (440/\% \text{NaOH}) + 4$ and % $\text{Zn} \approx (1240/\% \text{NaOH}) - 15$, resp. The soly. of ZnO at the b.p. of the solns., 121-315°, increases rapidly with the NaOH concn., passes through a max. at 56% NaOH (about 23.5% ZnO), and then decreases sharply. At the max., and beyond, the solid phase in equil. is represented by $4[\text{ZnO}.\text{Na}_2\text{O}].3\text{H}_2\text{O}$. The presence of NaCl decreases the soly. of ZnO .
1. Bepcew...

MT

URAZOV, G.G.; LOVCHIKOV, V.S.; LIPSHITS, B.M.

Zinc removal from lead by alkalization. TSvet.met.29 no.12:33-35 D
'56. (MLRA 10:2)
(Lead--Metallurgy)

PHASE I BOOK EXPLOITATION

406

Lipshits, Bella Moiseyevna
Suvorovskaya, Natal'ya Aleksandrovna; Titov Veleriy Ivanovich;
Brodsкая, Velentina Mikhaylovna; Vasil'yev, Pavel Ivanovich;
Lipshits, Bella Moiseyevna; and Eléntukh, Mariya Pavlovna

Tekhnicheskii analiz v tsvetnoy metallurgii (Technical Analysis
in Nonferrous Metallurgy) Moscow, Metallurgizdat, 1957.
567 p. 6,000 copies printed.

Reviewers: Troitskaya, M.I., Pomerantsev, I.N., Kozhukova, M.A.,
Candidates of Technical Sciences; Ed.: Vagina, N.S.; Ed.
of Publishing House: Kosolapova, E.F.; Tech Ed.:
Vaynshteyn, Ye. B.

PURPOSE: This is a textbook for use in technicums giving courses
in nonferrous metallurgy; it may also be used by those
performing chemical analysis at plant laboratories.

COVERAGE: The book describes widely used chemical and physico-
chemical methods of determining the constituents of nonferrous-
metal ores, of processed-ore products, of alloys, etc.

Card ~~1/42~~

Technical Analysis in Nonferrous Metallurgy 406

In addition, sections are included which are devoted to assaying, fuel analysis, water analysis, quality control in electrode production, and rational analysis. For authors of individual sections and chapters, see Table of Contents. There are 98 references, of which 85 are Soviet, 10 English, and 3 Czech.

TABLE OF CONTENTS:

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Technical analysis and its importance in quality control of metallurgical products	16
Methods of technical analysis	16
Selection of a representative sample	17
Principles of the separation of ions	20
Card 2/12	

URAZOV, G.G. [deceased]; LOVOHIKOV, V.S.; LIPSHITS, B.M.

Refining lead from arsenic, tin and antimony by liquid molten
alkali. Izv. vys. ucheb. zav.; tsvet. met. no.2:77-84 '58.
(MIRA 11:8)

1. Moskovskiy institut tsvetnykh metallov i zolota. Kafedra
tyazhelykh tsvetnykh metallov.
(Lead—Metallurgy)

AUTHOR: Urazov, G.G., (Deceased) SOV/149-58-4-13/26
Lovchikov, V.S.,
Lipshits, B.M.

TITLE: ~~Oxidation of Arsenic~~, Tin, and Antimony in Refining
Lead by Alkaline Melts (Okisleniye mysh'yaka, olova i
sur'my pri rafinirovanii svintsa shchelochnymi plavami)

PERIODICAL: Izvestiya Vysshikh Uchebnykh Zavedeniy, Tsvetnaya
Metallurgiya, 1958, Nr 4, pp 96-102 (USSR)

ABSTRACT: The results obtained by other workers (Ref.1-3) who
had studied kinetics of the reactions occurring when
fused NaOH and NaNO₃ are used for refining lead,
prompted the present Authors to investigate the
possibility of improving the efficiency of the refining
process by separate oxidation of the main impurities
(i.e. As, Sn and Sb) present in the crude metal. To
this end, the effect of various factors on the rate of
oxidation of these elements was studied in the
following manner: Oxidising mixtures of various
composition were added to impure lead melted in an
electrically heated iron crucible and maintained at

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SOV/149-58-4-13/26

Oxidation of Arsenic, Tin and Antimony in Refining Lead by
Alkaline Melts

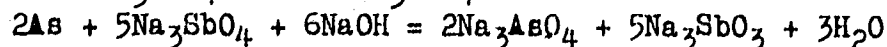
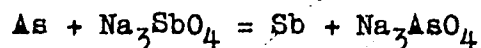
constant temperature and the whole was continuously stirred with a mechanical stirrer. Samples of metal and the salt were taken at regular intervals and chemically analysed. The results were plotted in the form of graphs showing how under various experimental conditions the impurity content in the refined metal changed with time. It was found that: In air, the rate of oxidation of As by NaOH is considerably higher at 450°C than at 400°C (Fig.1). In dry nitrogen this reaction occurs at approximately the same rate in the case of arsenic but neither Sn nor Sb are oxidised by NaOH under these conditions (Fig.2). In air, both Sn and Sb react with NaOH but at a much slower rate than As (Fig.3), which at 450°C is almost completely oxidised after 1 hour's treatment. Complete oxidation of arsenic in 5 minutes can be attained if instead of NaOH a mixture of 78% NaOH and 22% Na_3SbO_4 is employed (Fig.4). It can be seen from Fig.4 that in the course of the reaction with As, sodium antimonate is reduced

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to metallic Sb, so that oxidation of As takes place
according to the following reactions:



Similar reactions take place between Na_3SbO_4 and Sn (Fig.5) but in this case 16 hours are necessary completely to oxidise the impurity. According to the law of mass action, it is lead that is oxidised in the first place during the refining process. PbO reacts with NaOH forming sodium plumbite which reacts with As, Sn and Sb to yield metallic lead and corresponding arsenates. (Fig.6 shows the rate of oxidation of As, Sn and Sb by a melt consisting of 80% NaOH and 20% NaCl to which a quantity of PbO, 20% higher than that necessary to oxidise the impurities present in the metal, has been added: Arsenic is almost completely oxidised in

Card 3/5 10 minutes, while the content of metallic Sn and Sb is

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reduced in 3 hrs to 97 and 92% respectively). Although it would be possible to oxidise all these impurities with the aid of sodium plumbite, some lead oxide would remain after refining in the alkaline melt. For this reason it is necessary to oxidise the last traces of the impurities with NaNO_3 . (The results of experiments in which the alkaline melt consisted of 80% NaOH and 20% NaCl with a quantity of NaNO_3 theoretically necessary to oxidise As and Sb has been added, are shown on Fig.7.) In the course of the refining process the impurities present in the alkaline melt displace each other in the following order: As, Sn, Sb, Pb. However, it is not possible to obtain melt containing one of these elements only, owing to the fact that the displacement process does not proceed to completion. This has been shown not only by the results of laboratory experiments, but also by a large scale production test, the results of which are reproduced on Fig.8. It was concluded that separate recovery of

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Oxidation of Arsenic, Tin and Antimony in Refining Lead by
Alkaline Melts

the impurities under consideration may be expedient when the Sb content in the crude metal is several times higher than the content of the two other impurities (separately and jointly). In such a case, refining should be carried out in two stages: In the first stage As, Sn and a small proportion of Sb is recovered from the metal, while the bulk of the latter impurity is recovered in the second operation. There are 8 diagrams, 2 Soviet and 2 German references.

ASSOCIATION: Moskovskiy Institut Tsvetnykh Metallov i Zolota,
Kafedra Metallurgii Tyazhelykh Tsvetnykh Metallov
(Moscow Institute of Non-Ferrous Metals and Gold,
Chair for Metallurgy of Heavy Non-Ferrous Metals)

SUBMITTED: 15th April, 1958.

Card 5/5

LOVCHIKOV, V.S.; LIPSHITS, B.M.

Preparing antimony and tin from products of treating molten
alkali-refined lead. Izv. vys. ucheb. zav.; tsvet. met. 2
no.3:78-81 '59. (MIRA 12:9)

1. Moskovskiy institut tsvetnykh metallov i zolota, Kafedra
metallurgii tyazhelykh tsvetnykh metallov.
(Nonferrous metals--Metallurgy)

LOVCHIKOV, V.S.; LIPSHITS, B.M.

Production of tellurium concentrate fluxes produced by the
alkali refining of lead. Izv.vys.ucheb.sav.; tsvet.met. 2
no.6:93-98 '59. (MIRA 13:4)

1. Krasnoyarskiy institut tsvetnykh metallov. Kafedra
metallurgii tyazhelykh tsvetnykh metallov.
(Lead--Metallurgy) (Tellurium)

5(4)
 AUTHORS: Urazov, G. G. (Deceased), Lipshits, B. M., Lovchikov, V. S. SOV/78-4-2-29/40

TITLE: The Solubility Isotherms of the System $\text{Na}_2\text{O}-\text{H}_2\text{O}-\text{Sb}_2\text{O}_5$ at 25 and 75° (Izotermiy rastvorimosti sistemy $\text{Na}_2\text{O}-\text{H}_2\text{O}-\text{Sb}_2\text{O}_5$ pri 25 i 75°)

PERIODICAL: Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 2, pp 439-444 (USSR)

ABSTRACT: The solubility isotherms of the system $\text{Na}_2\text{O}-\text{H}_2\text{O}-\text{Sb}_2\text{O}_5$ were investigated for the first time at 25 and 75°. The synthesis of the compound $\text{NaSbO}_3 \cdot 3\text{H}_2\text{O}$ is given. The solubility of sodium antimoniate in water, depending on the temperature, was investigated and it was found that the solubility increases considerably upon a temperature rise. The isothermal solubility diagram of the system $\text{Na}_2\text{O}-\text{H}_2\text{O}-\text{Sb}_2\text{O}_5$ at 25 and 75° was drawn. The following salts crystallize in the system at 25°: $\text{NaSbO}_3 \cdot 3.5\text{H}_2\text{O}$; $\text{NaSbO}_3 \cdot 1.5\text{H}_2\text{O}$; $\text{Na}_3\text{SbO}_4 \cdot 6\text{H}_2\text{O}$ and Na_3SbO_4 . The crystallization zones of these salts depend on the concentration

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SOV/78-4-2-29/40
. The Solubility Isotherms of the System $\text{Na}_2\text{O}-\text{H}_2\text{O}-\text{Sb}_2\text{O}_5$ at 25 and 75°

of caustic soda. The salt $\text{NaSbO}_3 \cdot 1.5\text{H}_2\text{O}$ crystallizes difficultly. In this system the following salts crystallize at 75°: $\text{NaSbO}_3 \cdot 3\text{H}_2\text{O}$; $\text{Na}_3\text{SbO}_4 \cdot 6\text{H}_2\text{O}$ and Na_3SbO_4 . The existence of these salts also depends on the concentration of caustic soda. Upon investigation of the solubility of newly produced sodium antimoniate the salt showed comparatively great solubility in the concentration range of caustic soda from 40-49 weight % NaOH at 75°. There are 4 figures, 7 tables, and 5 references, 1 of which is Soviet.

ASSOCIATION: Moskovskiy institut tsvetnykh metallov i zolota im. M. I. Kalinina (Moscow Institute of Nonferrous Metals and Gold imeni M. I. Kalinin)

SUBMITTED: November 22, 1957

Card 2/2

5(2)

SOV/78-4-10-32/40

AUTHORS: Urazov, G. G. (Deceased), Lipshits, B. M., Lovchikov, V. S.

TITLE: Isotherms of Solubility in the System $\text{Na}_2\text{O} - \text{H}_2\text{O} - \text{SnO}_2$ at 25 and 75°

PERIODICAL: Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 10, pp 2380 - 2383 (USSR)

ABSTRACT: On refining lead by means of alkali an oxidation of ~~tin~~ occurs which passes over into the melt. In order to obtain the tin from the melt, the composition of the sodium salts of stannic acid and their solubility in sodium hydroxide must be known. To obtain these data, the system $\text{Na}_2\text{O} - \text{H}_2\text{O} - \text{SnO}_2$ was investigated.

The results are given in tables 1 and 2 and figures 1-3. At 25° and a concentration of NaOH between 10.95 - 36.9 wt% the salt $\text{Na}_2[\text{Sn}(\text{OH})_6]$ crystallizes. At 75° this salt crystallizes in the concentration range of NaOH between 20.74 - 45.88 wt%. At NaOH concentrations between 49.98 - 68.88 wt% the salt $\text{Na}_3[\text{Sn}(\text{OH})_7]\text{H}_2\text{O}$ is stable. At both temperatures the increasing NaOH-concentration involves a decreasing solubility of the stannate. The following practical conclusions are drawn from the afore-mentioned: The melts of the alkaline lead refining can be granulated in a solu-

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Isotherms of Solubility in the System $\text{Na}_2\text{O} - \text{H}_2\text{O} - \text{SnO}_2$ SOV/78-4-10-32/40
at 25 and 75°

tion with 35% NaOH. There the tin forms the well filterable salt $\text{Na}_2[\text{Sn}(\text{OH})_6]$. An increase of the sodium hydroxide concentration must be avoided because of the formation of a viscous pulp. The slime obtained by granulation of the melt can be filtered in hot state since the solubility of the sodium stannate in concentrated sodium lye is practically independent on temperature. The filter cake from sodium stannate must be washed with an alkaline solution, since it hydrolyzes with pure water. If it contains sodium antimonate, the sodium stannate can be separated from it by dissolving it by means of Ca-free water. There are 3 figures, 2 tables, and 4 references, 1 of which is Soviet.

SUBMITTED: July 11, 1958

Card 2/2

5(2)

SOV/80-32-5-11/52

AUTHORS: Urazov, G.G., Lipshits, B.M., Lovchikov, V.S.

TITLE: The Effect of Table Salt on the Solubility of Sodium Antimonate, Stannate and Arsenate in Aqueous Solutions of Caustic Soda

PERIODICAL: Zhurnal prikladnoy khimii, 1959, Vol 32, Nr 5, pp 995-997 (USSR)

ABSTRACT: The investigation of the oxidation process of arsenic, tin and antimony has shown that table salt, which is used in the refining of lead, does not take part in the reactions, decreases the chemical activity of alkali melts and increases the time needed for oxidation. Experiments are made here using caustic soda (purified of sodium carbonate), chemically pure sodium chloride and especially prepared sodium antimonate, arsenate and stannate. The solubility of sodium antimonate in water at 25°C is 0.10% at 75°C - 0.58%. Table salt decreases the solubility. The principal effect being obtained by caustic soda, the presence of table salt in the solution is unnecessary. At a content of 350 g/l NaOH the effect of sodium

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SOV/80-32-5-11/52

The Effect of Table Salt on the Solubility of Sodium Antimonate, Stannate and Arsenate in Aqueous Solutions of Caustic Soda

carbonate which reduces the solubility of sodium antimonate, is cancelled. Table salt decreases also the solubility of sodium stannate. At the mentioned content of NaOH the effect is not observed. The same effect is observed with sodium arsenate. The temperature is very important. On cooling sodium arsenate does not precipitate, but forms a crystalline structure. There are: 3 tables and 1 Soviet reference.

SUBMITTED: April 14, 1958

Card 2/2

URAZOV, G.G. [deceased]; LIPSHITS, B.M., LOVCHIKOV, V.S.

Solubility isotherm of the system $\text{Na}_2\text{O} - \text{H}_2\text{O} - \text{As}_2\text{O}_5$ at 75°
(concerning the process of the alkaline refining of lead).
Zhur. neorg. khim. 5 no.4:950-952 Ap '60. (MIRA 13:7)

1. Moskovskiy institut tsvetnykh metallov im. M.I. Kalinina.
(Sodium oxide) (Arsenic oxide) (Lead)

URAZOV, G.G. [deceased]; LIPSHITS, B.M.; LOVCHIKOV, V.S.

Mutual effect of sodium arsenate, stannate, and antimonate on their solubility in alkaline solutions. Zhur. neorg. khim. 5 no.11:2509-2511 N '60. (MIRA 13:11)

(Sodium arsenate)

(Sodium stannate)

(Sodium antimonate)

LIPSHITS, B.M.; SMIRNOVA, G.K.

Quantitative spot analysis of germanium with the use of phenylfluorone.
Zav.lab. 26 no.3:273-274 '60. (MIRA 13:6)

1. Institut tsvetnykh metallov i zolota im. M.I.Kalinina.
(Germanium--Analysis)
(Isocxanthenone)

LOVCHIKOV, V.S.; LIPSHITS, B.M.

Hydrometallurgical treatment of alkaline melts obtained as
a result of zinc removal from lead. TSvet.met. 33 no.1:76-77
Ja '60. (MIRA 13:6)
(Lead--Metallurgy) (Hydrometallurgy)

S/149/61/000/002/009/017
A006/A001

AUTHORS: Lovchikov, V.S., Lipshits, B.M., Obidina, L.A., Zubarev, Yu.V.
TITLE: On the Problem of Extracting Tellurium From Alkali Lead Refining Melts
PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy, Tsvetnaya metallurgiya, 1961, No. 2, pp. 97 - 101

TEXT: The hydrometallurgical processing of alkali lead refining melts is accompanied by the distribution of tellurium over all the products. Tellurium may be concentrated in sodium antimonate by precipitation from strong alkali solutions with antimony metal. (See tsvetnaya metallurgiya, # 6, p. 93, 1959). To determine optimum conditions of this process a series of experiments were performed. The initial solution contained 1.1 g/l Te; 350 g/l NaOH and 65 g/l NaCl. Tellurium was extracted from the solution with CY-2 (SU-2) grade antimony of the following grain sizes: - 3.2+1.5 mm, - 1.5+0.85 mm; - 0.85+0.42 mm and - 0.42+0.25 mm. When precipitating tellurium the theoretical amount of antimony of the aforementioned granulometric composition was consumed, and also its two-, four- and nine-fold excess in relation to the theoretical consumption. The experiments

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S/149/61/000/002/009/017
A006/A001 ✓

On the Problem of Extracting Tellurium From Alkali Lead Refining Melts


were performed in a glass container with a mechanical mixer into which 500 ml of the alkali solution were filled. The solution was heated to 95°C. An iron-grid basket containing antimony metal was placed into the hot solution and the mixer was switched on. The temperature and volume of the solution, and the rotation speed of the mixer were kept constant. Samples of the solution were subjected to chemical analysis, as to their tellurium content. The results show that higher consumption of antimony and smaller grain size raise the rate of separating tellurium out of the solution. It is recommended to conduct tellurium extraction from a strong alkaline solution at 95°C with a nine-fold excess of antimony over the theoretical amount at -0.82 ± 0.42 mm grain size for 3.5 hours. During reduction melting of sodium antimonate tellurium passes into the slag whose leaching out with water is accompanied by the formation of a solid residue containing over 3% Te. From this product Te may be leached out by an aqueous solution of sodium sulfide. To determine the optimum conditions of this process the authors studied the effect of temperature, the concentration of sodium sulfide in the initial solution, the liquid-solid ratio in the pulp and the time of leaching out. Leaching out of tellurium from the solid residue was made in a glass container with a

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A006/A001

On the Problem of Extracting Tellurium From Alkali Lead Refining Melts

mechanical mixer, using an aqueous solution of sodium sulfide and a solid residue, containing (in %): 3.2 Te; 14.0 Sb; 14.1 SiO₂; 7.51 CaO; 2.9 Fe₂O₃; 2.14 MgO and 0.18 Al₂O₃. The pulp volume and rotation speed of the mixer were maintained constant. The results obtained show that Te should be leached out from a solid residue by a solution containing 60 g/l Na₂S, for 5 hours at 95°C and 12:1 liquid-solid ratio in the initial pulp. This assures a 93% transition of Te into the solution. The solid residue (40%) contains (in %): 0.52 Te; 5.2 Sb; 29.7 SiO₂; 12.4 CaO; 4.1 Fe₂O₃; 3.8 MgO and 0.25 Al₂O₃. From the solution obtained tellurium was precipitated by sodium hydrosulfide (10 g per 1 g Te). Within 1.5 hours at 95°C, 95% Te in the form of metallic powder was extracted into the precipitate. The powder was extracted from the solution by filtrating the pulp. The dry powder contained 96% Te. After extracting tellurium a filtrate was obtained containing 32 g/l Na₂S and 20 g/l Na₂SO₃. The sodium hydrosulfide was removed from the solution with the aid of Ca(OH)₂. Optimum conditions for cleaning the sodium sulfide solution from sodium hydrosulfide were assured by using a 50% excess of calcium hydroxide in relation to the theoretical amount, and



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A006/A001

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On the Problem of Extracting Tellurium From Alkali Lead Refining Melts

stirring of the pulp for one hour at 95°C. The solution so obtained may be used for leaching out tellurium from new portions of solid residue. There are 9 figures and 1 Soviet reference.

ASSOCIATIONS: Krasnoyarskiy institut tsvetnykh metallov (Krasnoyarsk Institute of Nonferrous Metals), Kafedra metallurgii tyazhelykh tsvetnykh metallov (Department of Metallurgy of Heavy Non-Ferrous Metals)

SUBMITTED: May 18, 1960

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55300

27834
S/032/61/027/010/007/022
B110/B101AUTHORS: Lipshits, B. M., Smirnova, G. K., and Kulikov, F. S.

TITLE: Determination of iron in highly pure antimony

PERIODICAL: Zavodskaya laboratoriya, v. 27, no. 10, 1961, 1199 - 1200

TEXT: The determination of iron in Sb metal by means of thiocyanate is disturbed by the formation of a yellow precipitate. α , α' -dipyridyl forms a stable, soluble ferrodipyridyl complex ion, whose red color exactly obeys Lambert-Beer's law, and which is concentrated in a thin cresol layer.

1 - 5g of Sb metal was dissolved in a mixture of 5 parts of HCl 1:1 and 1 part of HNO_3 , (1:1), and evaporated to dryness at $<100^\circ\text{C}$ since otherwise

the iron volatilizes. The residue was dissolved in 50 ml solution of tartaric acid, NaCl, Na_2SO_3 , and NaOH purified from iron, (to form the soluble Sb complex), and boiled for 3 - 5 min. The pH should be 3 - 4. 2.0 ml α , α' -dipyridyl solution was added, filled up to 50 ml, and colorimetric measurement was conducted after 1 hr. When red coloring fails to appear, 15 - 20 ml Sb solution and 1 ml colorless cresol are

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Determination of iron in highly pure...

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B110/B101

filled in portions into a separating funnel, and shaken each time for 2 min. After extraction of the Fe complex, 0.1 μ g Fe in 5 g Sb may be visually determined by comparison with standard solutions. For plotting the calibration curve, standard solutions containing 0 - 5 μ g Fe are filled with 1 ml 10% hydroxylamine solution, 0.2% α , α' -dipyridyl solution, and 5 ml acetate buffer (0.5 ml glacial acetic acid and 0.3 g sodium acetate in 100 ml H₂O). The solutions are filled up to 25 ml with H₂O and

colorimetrically measured after 30 min on an Φ 3K-H-57 (FEK-N-57) with green light filter. Admixtures of Ni, Cd, As, Pb, Mn, Co, Bi, Ag, Pt, Au, Hg, Cu, Zn amounting to the 2 - 4 fold of the Fe content do not disturb. The acids used for dissolving Sb should be of special purity. 7.5 g NaCl, 15 g Na₂SO₃, 30 g tartaric acid, 10 - 11 g NaOH were dissolved in 150 ml

aqua dest., shaken, brought to pH = 4 - 5 by means of NaOH or HCl, and boiled for 3 - 5 min. 45 ml α , α' -dipyridyl solution was added and left standing for 18 - 20 hr under seal. 70 ml cresol was added in the separating funnel, and the Fe-free aqueous layer was filtered off. The solution was investigated for the presence of Fe by means of 0.5 ml α , α' -dipyridyl solution and color comparison with aqua dest. When red coloring failed X

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Determination of iron in highly pure...

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to appear, the solution was filled up to 600 ml with H₂O.

ASSOCIATION: Institut tsvetnykh metallov i zolota im. M. I. Kalinina
(Institute of Nonferrous Metals and Gold imeni M. I. Kalinin)

Card 3/3

LOVCHIKOV, V.S.; LIPSHITS, B.M.; OBIDINA, L.A.; ZUBAREV, Yu.V.

Extraction of tellurium from saturated lead leaching reagents.
Izv. vys. ucheb. zav.; tsvet. met. 4 no.2:97-101 '61.
(MIRA 14:6)

1. Krasnoyarskiy institut tsvetnykh metallov, kafedra metallurgii
tyazhelykh tsvetnykh metallov.
(Leaching)
(Tellurium--Metallurgy)

LOVCHIKOV, V.S.; LIPSHITS, B.M.

Hydrometallurgical treatment of the cake obtained in the alkali
refining of lead. Sbor. nauch. trud. GINTSVETMET no.33:43-50
'60. (MIRA 15:3)

(Lead--Metallurgy) (Hydrometallurgy)

LIPSHITS, B.M.; ANDREYCHUK, A.M.; AGAFONOVA, G.S.

Colorimetric determination of copper in metallic mercury. Zav.
lab. 30 no.9:1075 '64. (MIRA 18:3)

1. Moskovskiy institut stali i splavov.

BARYSHNIKOV, K.I.; BRISKIN, A.I.; VOROTYNTSEV, A.P.; GONCHAROV, P.I.;
DRUGOV, Yu.V.; LIPSHITS, L.A.; MOKEYEV, N.I.; NAZAROV, A.V.;
PETROV, L.P.; SERDYUK, D.S.; SNETANKIN, K.P.; CHERNYAVSKIY, A.A.;
ARTEM'YEV, S.G., red.; ZAKHAROVA, A.I., tekhn.red.

[Sanitary and chemical protection; pathology, clinical aspects,
and treatment of poisoning. Manual for students and physicians]
Sanitarno-khimicheskaya zashchita; patologiya, klinika i terapiya
porazhenii otravlyaiushchimi veshchestvami. Rukovodstvo dlia stu-
dentov i vrachei. Moskva, Gos.izd-vo med.lit-ry, 1959. 434 p.
(MIRA 13:6)

(CHEMICAL WARFARE---SAFETY MEASURES)

		1ST AND 2ND GRADERS		PROCESSES AND PROPERTIES INDEX		3RD AND 4TH GRADERS	
LIPSHITS, L A							IIIH
CA		Effect of early dosage with morphine in diphosgene poisoning. Z. M. Yavich, L. A. Lipshits, and N. F. Ochumkova. Farmakol. i Toksikol., 8, No. 1, 37-41 (1955). When given to dogs or rabbits within 30 min after diphosgene (I), morphine (II) decreases mortality in rats. II increases nerve excitability and is harmful. The favorable effect of II is attributed to decreased stimulation of the respiratory center and related parts of the central nervous system. In rabbits atropine (III) and O have a similar but less pronounced effect on respiration, but they increase the pulse rate whereas II lowers it. Hemoglobin count in rabbits is raised by III, lowered by I and II; body temp. is lowered slightly by O, more by III and II; (in the first 3 hrs.) still more by II. It seems probable that prompt dosage with II would be helpful to patients poisoned with I.					
Juhon E. Smith							
METALLURGICAL LITERATURE CLASSIFICATION		RESEARCH NUMBER					
ASB-SLA		SELECT ONE ONLY					
FROM STUDYING		ALSO SEE					
DATE		NO.					
AV		C					
MAY		D					
1955		E					
10		F					
11		G					
12		H					
13		I					
14		J					
15		K					
16		L					
17		M					
18		N					
19		O					
20		P					
21		Q					
22		R					
23		S					
24		T					
25		U					
26		V					
27		W					
28		X					
29		Y					
30		Z					

LIFSHITS, M. (Saratov)

Evaluation of the exterior. Voen. zhnan. 40 no.8:46-47 S 162.
(REF 17:12)

LIPSHITS, N. V.

Looms

Central woof fork on the machines At-175 SH Tekst. prom. no. 5, 1952.

Monthly List of Russian Accessions, Library of Congress, August 1952. Unclassified.

LIPSHITS, N.V.

LIPSHITS, N.V.

Possibility for reducing the consumption of yarns in weaving.

Tekst. prom. 14 no.5:48 My '54.

(MIRA 7:6)

(Weaving)

LIPSHITS, N.V.

The mechanism of double-action weft forks and the reverse motion
of the batten on BI-N looms. Tekst.prom. 15 no.2:27-30 F '55.
(Looms) (MLRA 8:3)

LIPSHITS, N.V.

Type of weft safety device on wide automatic looms. Tekst.prom.
15 no.9:18-21 S '55. (MIRA 8:11)
(Looms)

LIPSHITS, Naum Veniaminovich; BITUNOV, Ye.I., retsenzent; SOKOLOVA, V.Ye.,
redaktor; BEL'CHIKOVA, Yu.S., tekhnicheskiiy redaktor

[Organisation and maintenance of AT-175sh automatic loom]
Ustroistvo i obsluzhivanie avtomaticheskikh tkatskikh stankov
AT-175sh. Moskva, Gos. nauchno-tekhn. izd-vo Ministerstva legkoi
promyshl. SSSR, 1956. 137 p. (MLRA 9:9)
(Looms)

LIPSHITS, N. V. Cand Tech Sci -- (diss) ^{"Study of"} ~~Research~~ on guard devices on automatic
looms in the manufacture of woolen fabrics and the normalization of their work"
Mos, 1957. 14 pp 22 cm. (in ^{of} Higher Ed USSR. Mos Textile Institute), 100 copies
(H, 20-57, 64)

LIPSHITS, N. V.

LIPATENKOV, Ivan Vasil'yevich; KAPRALOV, Mikhail Karpovich; BITUNOV, Yevgeniy Ivanovich; VAKUROV, Konstantin Viktorovich; KUZOVSKIN, Konstantin Sergeyevich; PAVLOV, Leonid Vasil'yevich; KLOCHKOV, Ivan Nikitich; ZHITS, Margoliya Isayevna; KHROMOV, Vasil'y Vasil'yevich; LIPSHITS, N.V., redaktor; KOPILVICH, Ye.I., redaktor; DMITRIYEVA, N.I., tekhnicheskii redaktor

[Assembling and adjusting machinery of looms with picker sticks;
work practices of foremen and assistants in the Monin worsted mills]
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(over)

Vascular Permeability

bility appeared to have had no effect on leucocyte emigration. No parallelism was discerned between the intensity of ATP chemotaxis and changes in vascular permeability. Combined urethan and veronal narcotic sleep was produced in rabbits and the effect of ATP injection was studied 8-24 hrs. from the onset of narcosis. Under these conditions the vascular permeability was enhanced, and the deposition of the trypan blue in the injected areas was vastly increased. Expts. with rabbits lead to the conclusion that in narcotic sleep ATP does not manifest any noteworthy vascular permeability changes or chemotaxis. The action of ATP on vascular permeability is dependent upon the functional state of the higher regions of the nervous system and reflex processes lie at the base of ATP action on vascular permeability and the emigration of leucocytes. B. S. Levine

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